



uromodulin-associated kidney disease

Uromodulin-associated kidney disease is an inherited condition that affects the kidneys. The signs and symptoms of this condition vary, even among members of the same family.

Many individuals with uromodulin-associated kidney disease develop high blood levels of a waste product called uric acid. Normally, the kidneys remove uric acid from the blood and transfer it to urine. In this condition, the kidneys are unable to remove uric acid from the blood effectively. A buildup of uric acid can cause gout, which is a form of arthritis resulting from uric acid crystals in the joints. The signs and symptoms of gout may appear as early as a person's teens in uromodulin-associated kidney disease.

Uromodulin-associated kidney disease causes slowly progressive kidney disease, with the signs and symptoms usually beginning during the teenage years. The kidneys become less able to filter fluids and waste products from the body as this condition progresses, resulting in kidney failure. Individuals with uromodulin-associated kidney disease typically require either dialysis to remove wastes from the blood or a kidney transplant between the ages of 30 and 70. Occasionally, affected individuals are found to have small kidneys or kidney cysts (medullary cysts).

Frequency

The prevalence of uromodulin-associated kidney disease is unknown. It accounts for fewer than 1 percent of cases of kidney disease.

Genetic Changes

Mutations in the *UMOD* gene cause uromodulin-associated kidney disease. This gene provides instructions for making the uromodulin protein, which is produced by the kidneys and then excreted from the body in urine. The function of uromodulin remains unclear, although it is known to be the most abundant protein in the urine of healthy individuals. Researchers have suggested that uromodulin may protect against urinary tract infections. It may also help control the amount of water in urine.

Most mutations in the *UMOD* gene change single protein building blocks (amino acids) used to make uromodulin. These mutations alter the structure of the protein, preventing its release from kidney cells. Abnormal buildup of uromodulin may trigger the self-destruction (apoptosis) of cells in the kidneys, causing progressive kidney disease.

Inheritance Pattern

This condition is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder.

Other Names for This Condition

- Familial gout-kidney disease
- Familial gouty nephropathy
- Familial juvenile hyperuricemic nephropathy
- FJHN
- MCKD2
- Medullary cystic kidney disease type 2
- UMAK
- UMOD-related kidney disease
- Uromodulin storage disease

Diagnosis & Management

Genetic Testing

- Genetic Testing Registry: Familial juvenile gout
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0268113/>
- Genetic Testing Registry: Glomerulocystic kidney disease with hyperuricemia and isosthenuria
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1835934/>
- Genetic Testing Registry: Medullary cystic kidney disease 2
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1859040/>

Other Diagnosis and Management Resources

- GeneReview: Autosomal Dominant Tubulointerstitial Kidney Disease, UMOD-Related
<https://www.ncbi.nlm.nih.gov/books/NBK1356>

General Information from MedlinePlus

- Diagnostic Tests
<https://medlineplus.gov/diagnostictests.html>
- Drug Therapy
<https://medlineplus.gov/drugtherapy.html>

- Genetic Counseling
<https://medlineplus.gov/geneticcounseling.html>
- Palliative Care
<https://medlineplus.gov/palliativecare.html>
- Surgery and Rehabilitation
<https://medlineplus.gov/surgeryandrehabilitation.html>

Additional Information & Resources

MedlinePlus

- Health Topic: Gout
<https://medlineplus.gov/gout.html>
- Health Topic: Kidney Diseases
<https://medlineplus.gov/kidneydiseases.html>
- Health Topic: Kidney Failure
<https://medlineplus.gov/kidneyfailure.html>

Genetic and Rare Diseases Information Center

- Autosomal dominant medullary cystic kidney disease with hyperuricemia
<https://rarediseases.info.nih.gov/diseases/10679/autosomal-dominant-medullary-cystic-kidney-disease-with-hyperuricemia>

Additional NIH Resources

- National Institute of Arthritis and Musculoskeletal and Skin Diseases: Gout
https://www.niams.nih.gov/Health_Info/Gout/
- National Institute of Diabetes and Digestive and Kidney Diseases: The Kidneys and How They Work
<https://www.niddk.nih.gov/health-information/kidney-disease/kidneys-how-they-work>

Educational Resources

- Disease InfoSearch: Medullary cystic kidney disease 2
<http://www.diseaseinfosearch.org/Medullary+cystic+kidney+disease+2/4550>
- Merck Manual Home Edition: Gout
<http://www.merckmanuals.com/home/bone-joint-and-muscle-disorders/gout-and-pseudogout/gout>

- Orphanet: Autosomal dominant medullary cystic kidney disease with or without hyperuricemia
http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=34149
- Wake Forest University Baptist Medical Center
<http://www.wakehealth.edu/nephrology/gout/>

Patient Support and Advocacy Resources

- National Kidney Foundation
<https://www.kidney.org/>

GeneReviews

- Autosomal Dominant Tubulointerstitial Kidney Disease, UMOD-Related
<https://www.ncbi.nlm.nih.gov/books/NBK1356>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28uromodulin-associated+kidney+disease%5BTIAB%5D%29+OR+%28familial+juvenile+hyperuricemic+nephropathy%5BTIAB%5D%29+OR+%28fjhn%5BTIAB%5D%29+OR+%28mckd2%5BTIAB%5D%29+OR+%28uromodulin+storage+diseases%5BTIAB%5D%29+OR+%28medullary+cystic+kidney+disease+type+2%5BTIAB%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

OMIM

- GLOMERULOCYSTIC KIDNEY DISEASE WITH HYPERURICEMIA AND ISOSTHENURIA
<http://omim.org/entry/609886>
- HYPERURICEMIC NEPHROPATHY, FAMILIAL JUVENILE, 1
<http://omim.org/entry/162000>
- MEDULLARY CYSTIC KIDNEY DISEASE 2
<http://omim.org/entry/603860>

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